

AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

Claim 1(Currently Amended): An antigenic composition comprising

(a) at least one antigen from a pathogenic organism selected from the

group consisting of a bacterium, a virus, a fungus and a parasite; and

(b) an effective adjuvanting amount of a mutant cholera holotoxin, wherein the mutant holotoxin has reduced toxicity compared to a wild-type cholera holotoxin, and has a substitution an amino acid which replaces the deleted glutamic acid which naturally occurs at position 29 of the mature A subunit of the wild-type cholera holotoxin, with an wherein said amino acid is other than aspartic acid, and wherein said mutant holotoxin enhances the immune response in a vertebrate host to said antigen.

Claim 2 (Previously Presented): The antigenic composition of Claim 1 wherein the antigenic composition comprises more than one antigen of (a).

Claim 3(Previously Presented): An antigenic composition comprising

(a) at least one antigen from a pathogenic organism selected from the group consisting of a bacterium, a virus, a fungus and a parasite; and

(b) an effective adjuvanting amount of a mutant cholera holotoxin, wherein the mutant holotoxin has reduced toxicity compared to wild-type cholera holotoxin, and has a histidine which replaces the glutamic acid which naturally occurs at position 29 of the A subunit of the wild-type cholera holotoxin and wherein said mutant holotoxin enhances the immune response in a vertebrate host to said antigen.

Claim 4(Previously Presented): The antigenic composition of Claim 1 wherein the antigen is selected from the group consisting of the *Haemophilus influenzae* P4 outer membrane protein, the *Haemophilus influenzae* P6 outer membrane protein, the

Haemophilus influenzae adherence and penetration protein (Hap_s), the *Helicobacter pylori* urease protein, the *Neisseria meningitidis* Group B recombinant class 1 pilin (rpilin), the *Neisseria meningitidis* Group B class 1 outer membrane protein (PorA), the respiratory syncytial virus fusion protein, a rotavirus virus-like particle and the herpes simplex virus (HSV) type 2 glycoprotein D (gD2).

Claim 5(Previously Presented): The antigenic composition of Claim 4 wherein the antigen is selected from the group consisting of the *Haemophilus influenzae* P4 outer membrane protein, the *Haemophilus influenzae* P6 outer membrane protein, the *Haemophilus influenzae* Hap_s protein, and any combination thereof.

Claim 6(Previously Presented): The antigenic composition of Claim 4 wherein the antigen is the *Helicobacter pylori* urease protein.

Claim 7(Previously Presented): The antigenic composition of Claim 4 the antigen is selected from the group consisting of the *Neisseria meningitidis* rpilin, *Neisseria meningitidis* PorA protein and any combination thereof.

Claim 8(Previously Presented): The antigenic composition of Claim 4 wherein the antigen is the respiratory syncytial virus fusion protein.

Claim 9(Previously Presented): The antigenic composition of Claim 4 wherein the antigen is a rotavirus virus-like particle.

Claim 10(Original): The antigenic composition of Claim 9 wherein the virus-like particle is a rotavirus 2/6-virus-like particle.

Claim 11(Previously Presented): The antigenic composition of Claim 4 wherein the antigen is HSV gD2.

Claim 12(Canceled)

Claim 13(Original): The antigenic composition of Claim 1 wherein the antigenic composition further comprises a diluent or carrier.

Claim 14(Original): The antigenic composition of Claim 1 which further comprises a second adjuvant in addition to the mutant cholera holotoxin.

Claim 15(Currently Amended): The antigenic composition of Claim 1, wherein at least one additional mutation is made to the A subunit of the mutant cholera holotoxin at a position other than said wild-type amino acid position 29, ~~wherein said mutant holotoxin with said additional mutation enhances the immune response in a vertebrate host to said antigen~~.

Claim 16(Previously Presented): The antigenic composition of Claim 15 wherein the at least one additional mutation is made as a substitution for a naturally-occurring amino acid at an amino acid position of wild-type cholera holotoxin selected from the group consisting of the arginine at amino acid 7, the aspartic acid at position 9, the arginine at position 11, the histidine at position 44, the valine at position 53, the arginine at position 54, the serine at position 61, the serine at position 63, the histidine at position 70, the valine at position 97, the tyrosine at position 104, the proline at position 106, the histidine at position 107, the serine at position 109, the glutamic acid at position 100, the glutamic acid at position 112, the serine at position 114, the tryptophan at position 127, the arginine at position 146 and the arginine at position 192.

Claim 17(Previously Presented): A method for increasing the ability of an antigenic composition containing at least one antigen from a pathogenic organism selected from the group consisting of a bacterium, a virus, a fungus or a parasite to elicit the immune response of a vertebrate host, which comprises administering to said host an antigenic composition of Claim 1.

Claims 18-27(Canceled)

Claim 28 (Previously Presented): The method of Claim 17 wherein the antigenic composition comprises more than one antigen.

Claim 29(Previously Presented): The method of Claim 17 wherein the amino acid substituted at wild-type position 29 is histidine.

Claim 30(Previously Presented): The method of Claim 17 wherein the antigen is selected from the group consisting of the *Haemophilus influenzae* P4 outer membrane protein, the *Haemophilus influenzae* P6 outer membrane protein, the *Haemophilus influenzae* Hap_s protein, the *Helicobacter pylori* urease protein, the *Neisseria meningitidis* rpilin, the *Neisseria meningitidis* PorA protein, the respiratory syncytial virus fusion protein, a rotavirus, virus-like particle and HSV gD2.

Claim 31(Previously Presented): The method of Claim 30 wherein at least one antigen is selected from the group consisting of the *Haemophilus influenzae* P4 outer membrane protein, the *Haemophilus influenzae* P6 outer membrane protein, the *Haemophilus influenzae* Hap_s protein, and any combination thereof.

Claim 32(Previously Presented): The method of Claim 30 wherein the antigen is the *Helicobacter pylori* urease protein.

Claim 33(Previously Presented): The method of Claim 30 wherein at least one antigen is selected from the group consisting of the *Neisseria meningitidis* rpilin, *Neisseria meningitidis* PorA protein and any combination thereof.

Claim 34(Previously Presented): The method of Claim 30 wherein the antigen is the respiratory syncytial virus fusion protein.

Claim 35(Previously Presented): The method of Claim 30 wherein the antigen is a rotavirus virus-like particle.

Claim 36(Original): The method of Claim 35 wherein the virus-like particle is a rotavirus 2/6-virus-like particle.

Claim 37(Previously Presented): The method of Claim 30 wherein the antigen is HSV gD2.

Claim 38(Canceled)

Claim 39(Original): The method of Claim 17 wherein the antigenic composition further comprises a diluent or carrier.

Claim 40(Original): The method of Claim 17 wherein the antigenic composition further comprises a second adjuvant in addition to the mutant cholera holotoxin.

Claim 41(Previously Presented): The method of Claim 17 wherein at least one additional mutation is made to the A subunit of the mutant cholera holotoxin at a position other than said wild-type amino acid position 29, wherein said mutant holotoxin with said additional mutation enhances the immune response in a vertebrate host to said antigen.

Claim 42(Previously Presented): The method of Claim 41 wherein the at least one additional mutation is made as a substitution for a naturally-occurring amino acid of wild-type cholera holotoxin selected from the group consisting of the arginine at amino acid 7, the aspartic acid at position 9, the arginine at position 11, the histidine at position 44, the valine at position 53, the arginine at position 54, the serine at position 61, the serine at position 63, the histidine at position 70, the valine at position 97, the tyrosine at position 104, the proline at position 106, the histidine at position 107, the serine at position 109, the glutamic acid at position 100, the glutamic acid at position 112, the serine at position 114, the tryptophan at position 127, the arginine at position 146 and the arginine at position 192.

Claim 43(Previously Presented): A method of preparing an antigenic composition comprising combining

(a) at least one antigen from a pathogenic organism selected from the group consisting of a bacterium, a virus, a fungus and a parasite; and

(b) an effective adjuvanting amount of a mutant cholera holotoxin, wherein the mutant holotoxin has reduced toxicity compared to wild-type cholera holotoxin and has a substitution which replaces the glutamic acid which naturally occurs at position 29 of the A subunit of the wild-type cholera holotoxin with an amino acid other than aspartic acid, and wherein said mutant holotoxin enhances the immune response in a vertebrate host to said antigen.

Claim 44(Previously Presented): A method for increasing the ability of an antigenic composition containing at least one antigen from a pathogenic organism selected from the group consisting of a bacterium, a virus, a fungus or a parasite to elicit the immune response of a vertebrate host, which comprises administering to said host an antigenic composition of Claim 3.